Application No. 10/532,398 Docket No.: 0365-0627PUS1 Response Dated July 7, 2008

Reply to Office Action of March 5, 2008

AMENDMENTS TO THE CLAIMS

1. (Currently Amended) Process A process for manufacturing transglycosylation

products, according to which process comprising the steps of:

- reacting a starch derivative ester or starch ether is reacted at acidic conditions with an

alkanol containing 1 [[-]] to 6 hydroxyl groups in the presence of an acidic catalyst in a

transglycosylation reaction to form a reaction mixture, wherein said acid catalyst comprises

phosphorous, and

- the reaction recovering a transglycosylation product is recovered as such, or [[it]]

subjecting the transglycosylation product is subjected to further processing,

characterized in that

- the transglycosylation reaction between the starch derivatives and the alkanol is carried

out performed in a reactive extrusion process essentially without any medium, and

- the reaction mixture is conducted through an extrusion device via at least two separately

adjustable heating zones, thereby providing control of heat introduced externally into the

reaction mixture,

wherein the acidic catalyst is a phosphorus-containing acid selected from at least one of

 $\underline{\text{the group consisting of: phosphoric acid, $H_3PO_4$, hypophosphorous acid, $H_3PO_2$, and}\\$ 

phosphorous acid, H<sub>3</sub>PO<sub>3</sub>, and

wherein the catalyst is allowed to chemically bond with the transglycosylation product.

2-4. (Cancelled)

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5. (Currently Amended) The process according to claim 1, characterized in that wherein

the extrusion temperature is within the range of approx. approximately 105 to 200 °C, preferably

within the range of approx. 110 to 190 °C.

6. (Currently Amended) The process according to claim 1, characterized in that, wherein

prior to performing the transglycosylation reaction, the alkanol and the acidic substance catalyst

are mixed together to form a first reaction mixture, [[and]] thereby producing an aerosol is

produced from this mixture, [[and]] wherein the acrosol is added to the starch derivative at a

dose corresponding to [[the]] a desired molar mass of the [[end]] transglycolsytion product to

produce a pre-mixture.

7. (Currently Amended) The process according to claim 6, characterized in that wherein

the amount of the alkanol is approx. approximately 0.01 to 20 weight-%, preferably 0.1 to 10

weight %, of the mass of the starch derivative ester or the starch ether, and wherein the amount

of the alkanol is approx. approximately 0.0005 to approx. approximately 5 mole-%, preferably

approx. 0.002 to approx. 2.0 mole-%, in particularly approx 0.015 to 0.3 mole %, of the amount

of the starch derivative used ester or the starch ether.

8. (Currently Amended) The process according to claim 6 or 7, characterized in that

wherein the alkanol and the acidic substance catalyst are [[fed]] supplied in aerosol form [[into]]

 $\underline{to}$  a fluidised-bed type of a mixing device, in which they wherein the alkanol and acidic catalyst

are mixed with a powdery starch derivative to produce the pre-mixture.

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9. (Currently Amended) The process according to claim 6, eharacterized in that wherein

the concentrations of the alkanol and the acidic substance catalyst and of any liquid chemicals

are selected such that the total amount of liquid will be is less than 30 %, preferably approx. 5 to

25 %, of the dry matter content of the pre-mixture containing the starch derivative.

10. (Currently Amended) The process according to claim 5, characterized in that wherein

the reaction mixture is compacted and, where necessary, granulated vielding a compacted pre-

mixture, prior to feeding it into supplying the reaction mixture to the extrusion device.

11. (Currently Amended) The process according to claim 10, characterized in that

wherein the compacted pre-mixture is fed into supplied to the extruder extrusion device to serve

as the reaction mixture, wherein the extruder extrusion device being of is either [[the]] a 1- or 2-

screw type extrusion device.

12. (Currently Amended) The process according to claim 1, characterized in that wherein

the starch derivative ester or starch ether comprises a product manufactured from native starch

by means of oxidizing, hydrolyzing, cross-linking, cationizing, grafting, etherification or

esterification.

13. (Cancelled)

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14. (Currently Amended) The process according to claim 1, characterized in that wherein

the alkanol [[used]] is a lower alkanol with 1 to 6 carbon atoms [[,]] and 1 to 5, preferably 1 to 3,

hydroxyl groups.

15. (Currently Amended) The process according to claim 14, characterized in that

wherein the alkanol is selected from at least one of the group consisting of; methanol, ethanol, n-

propanol, isopropanol, n-butanol and see, sec butanol, a substituted lower alcohol, e.g. metoxy

methoxy ethanol, etoxy ethanol, metoxy methoxy methanol, or etoxy ethoxy methanol, or

an alcohol containing 2 or 3 hydroxyl groups, e.g. ethylene glycol, propylene glycol [[or]] and

glycerol.

16. (Currently Amended) The process according to claim 1, characterized in that wherein

the acid catalyst used is a strong mineral acid, such as is selected from at least one of the group

consisting of: sulphuric acid, hydrochloric acid, nitric acid, strong-organic acid, such as

paratoluene sulphonic acid, methane sulphonic acid, benzene sulphonic acid [[or]],

trifluoromethane sulphonic acid or mono or polyalkylated aryl mono or polysulphonic acid,

such as, xylene sulphonic acid [[or]], cumene sulphonic acid [[or]], and dodecyl benzene

sulphonic acid, or an acidic ion exchange resin.

17-21. (Cancelled)

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22. (New) The process according to claim 1, wherein the extrusion temperature is within

the range of approximately 110 to 190 °C.

23. (New) The process according to claim 6, wherein the amount of the alkanol is

approximately 0.1 to 10 weight-% of the mass of the starch ester or the starch ether.

24. (New) The process according to claim 6, wherein the amount of the alkanol is

approximately 0.002 to approximately 2.0 mole-% of the amount of the starch ester or the starch

ether.

25. (New) The process according to claim 6, wherein the amount of the alkanol is

approximately 0.015 to 0.3 mole-% of the amount of the starch ester or the starch ether.

26. (New) The process according to claim 6, wherein the concentrations of the alkanol

and the acidic catalyst and of any liquid chemicals are selected such that the total amount of

liquid is approximately 5 to 25 % wt-% of the dry matter content in the pre-mixture.

27. (New) The process according to claim 5, wherein the reaction mixture is compacted

and granulated prior to feeding it into the extrusion device.

28. (New) The process according to claim 1, wherein the alkanol is a lower alkanol with

1 to 6 carbon atoms and 1 to 3 hydroxyl groups.

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29. (New) The process according to claim 14, wherein the alkanol is selected from the

group consisting of: a substituted lower alcohol, an alcohol containing two hydroxyl groups, and

an alcohol containing three hydroxyl groups.

30. (New) The process according to claim 1, wherein the acid catalyst is selected from at

least one of the group consisting of: a strong mineral acid, mono-polyalkylated aryl

monosulphonic acid, mono-polyalkylated polysulphonic acid, polyalkylated aryl mono-sulphonic

acid, polyalkylated aryl polysulphonic acid, and an acidic ion exchange resin.

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